

Phosphate glass fibrous scaffolds: tailoring of the properties and improvement of the bioactivity through the incorporation of mesoporous glasses

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## Introduction

Synthetic bone **scaffolds** are proposed as an alternative to the use of bone grafting technique for bone regeneration. Porous scaffold obtained from cutting glass fibres and randomly arranging into a mould, shows the **open porosity** necessary for tissue ingrowth and vascularization. Moreover the use of a **resorbable glass** and **mesoporous bioactive particles** (*i.e.* specific surface area up to 800 m<sup>2</sup>/g, adjustable pore size between 2 and 50 nm, large pore volume [1]) allows to obtain a 3D structure in which the newly regenerated bone substitutes the synthetic material.

## Materials and methods

### Phosphate glass fibers

#### TiPS<sub>2.5</sub>

Fibres of a TiO<sub>2</sub>-containing phosphate glass fabricated following the preform drawing approach [2].

#### CEL2

Dense silica-based bioactive glass (45SiO<sub>2</sub>, 3P<sub>2</sub>O<sub>5</sub>, 26CaO, 7MgO, 15Na<sub>2</sub>O, 4K<sub>2</sub>O mol.%) produced by melt quenching technique [3].

#### SD\_MBG

Micro-sized mesoporous glass based on SiO<sub>2</sub>-CaO (80SiO<sub>2</sub>, 20 CaO mol.%) system produced by an aerosol-assisted spray-drying technique [4].

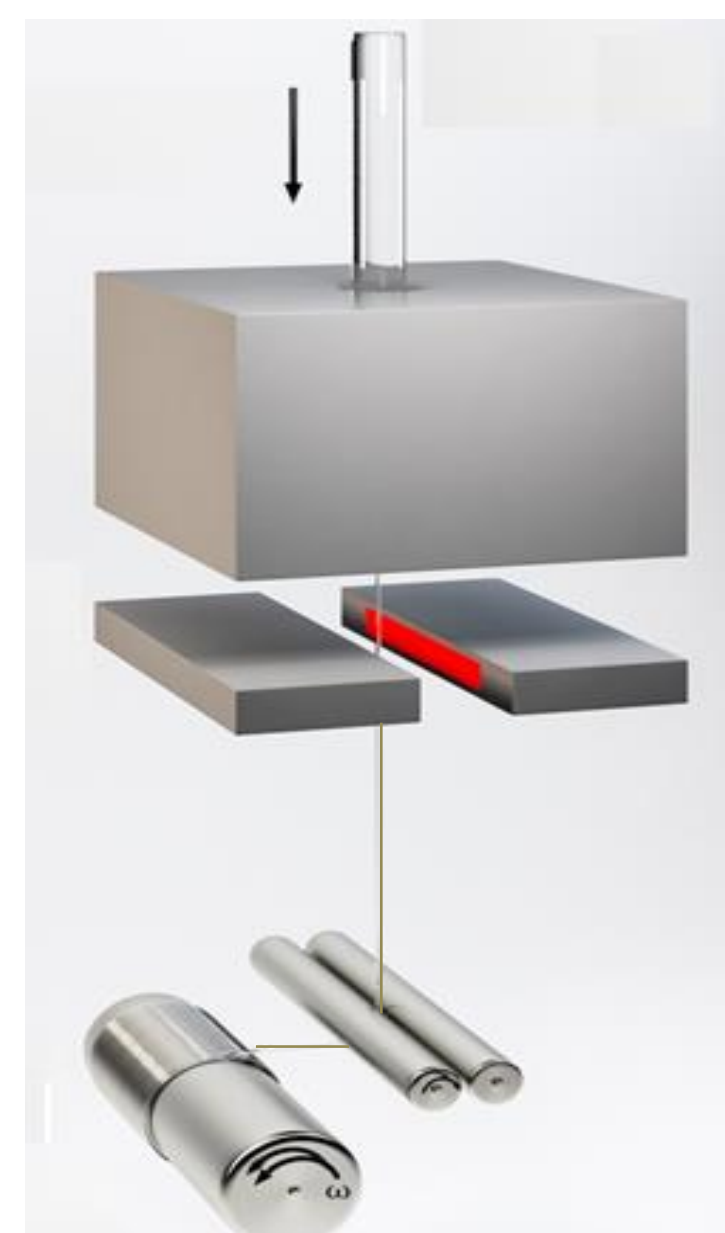
#### Cu\_BGN2%

Cu-containing mesoporous glass nanoparticles (85SiO<sub>2</sub>, 13 CaO, 2CuO mol.%) synthesized by an ultra-sound assisted sol-gel method.

### Scaffold preparation

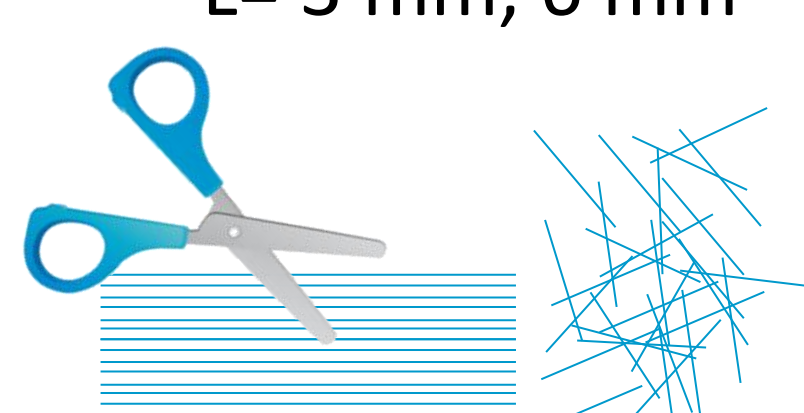
#### FIBRE DRAWING

Ø 110 µm



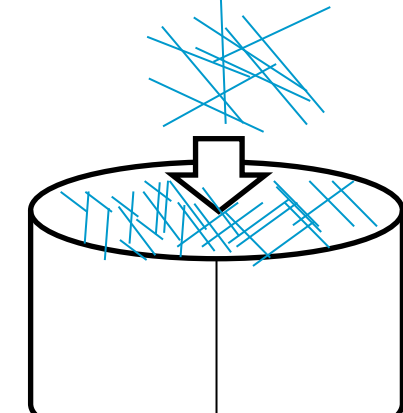
#### CUTTING

L = 3 mm, 6 mm



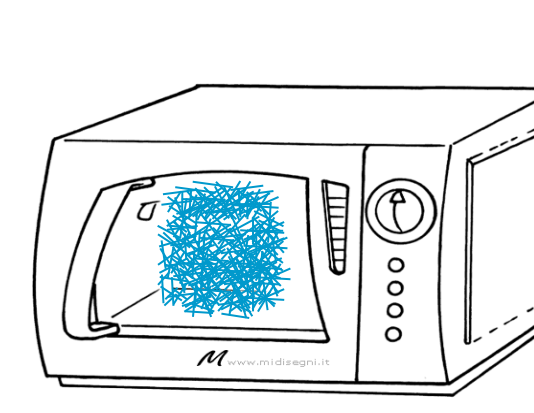
#### SHAPING

The structure shape is maintained after mould removal



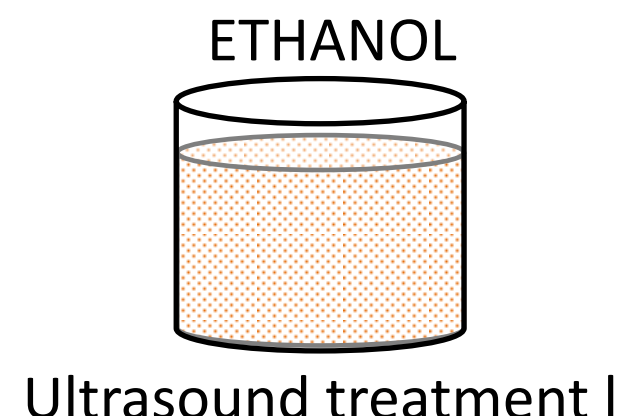
(D 13mm, h 12 mm)

#### SINTERING



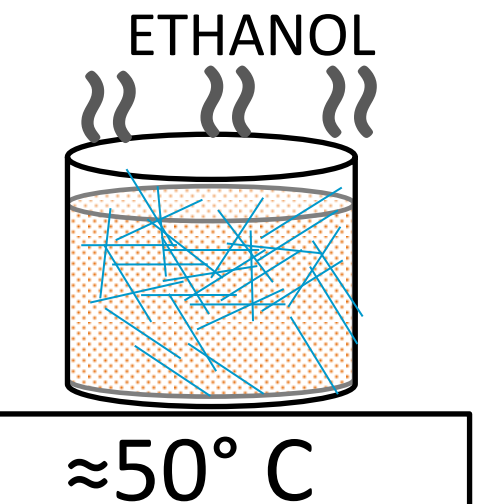
### ADDITIONAL STEP: INTRODUCTION OF THE BIOACTIVE POWDER

SUSPENSION OF BIOACTIVE POWDER IN ETHANOL



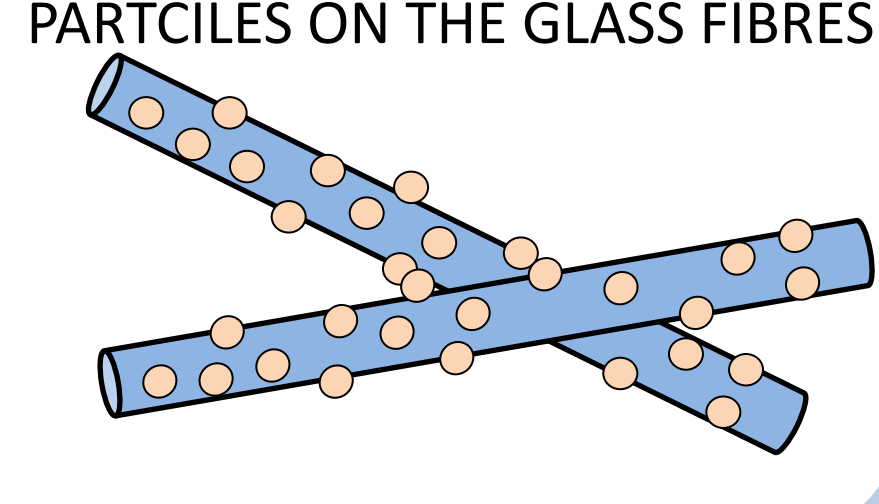
Ultrasound treatment I

EVAPORATION OF ETHANOL



≈50° C

DEPOSITION OF THE BIOACTIVE PARTICLES ON THE GLASS FIBRES



### Mesoporous Powder characterization

Structural analysis: N<sub>2</sub> adsorption/desorption technique

Morphological analysis: FESEM

### Scaffold characterization

Morphological analysis: FESEM

Inner structure: Micro-CT

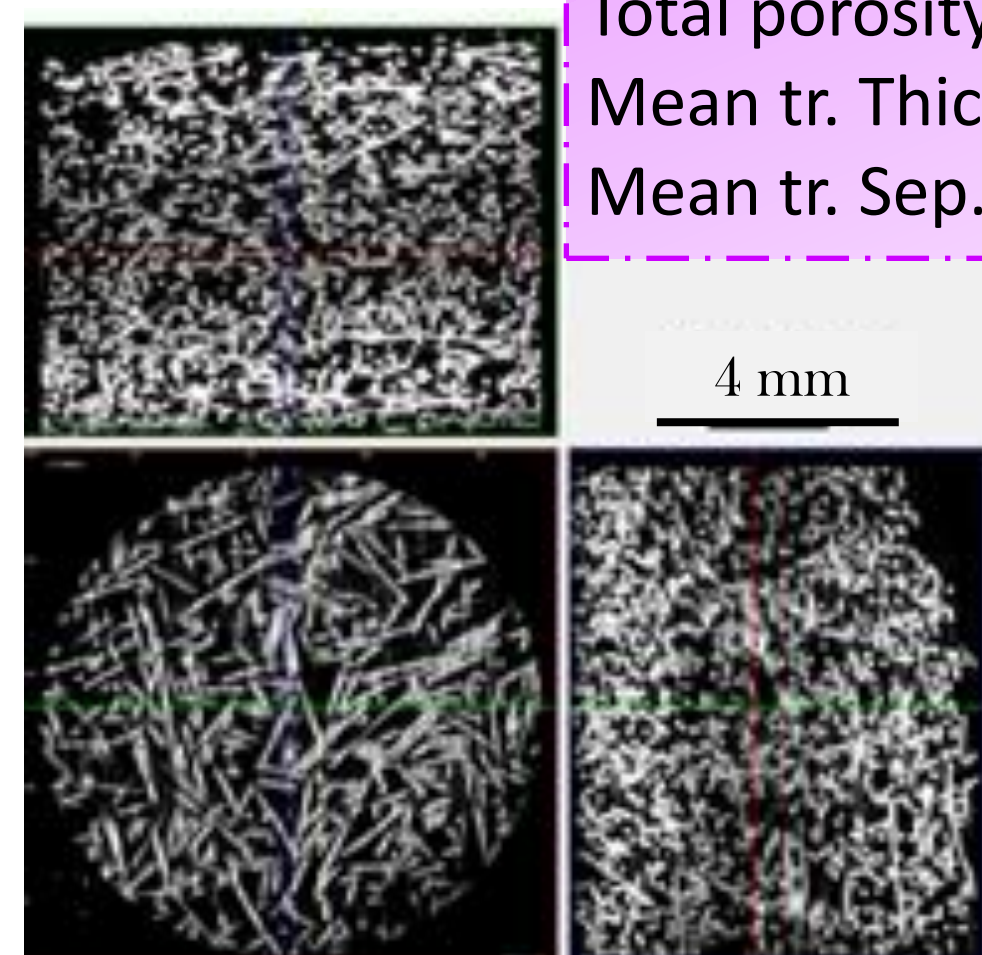
Scaffold bioactivity: SBF soaking test

## Results and discussion

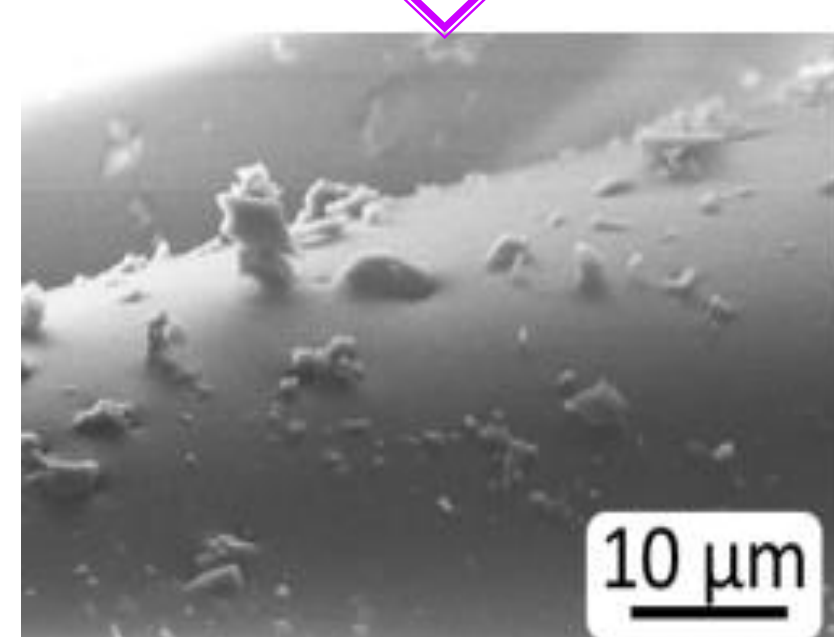
### S110\_CEL2

- 3 wt.% bioactive powder
- Fibres: 3 mm

Total porosity: 58%  
Mean tr. Thick :111 µm  
Mean tr. Sep.: 162 µm

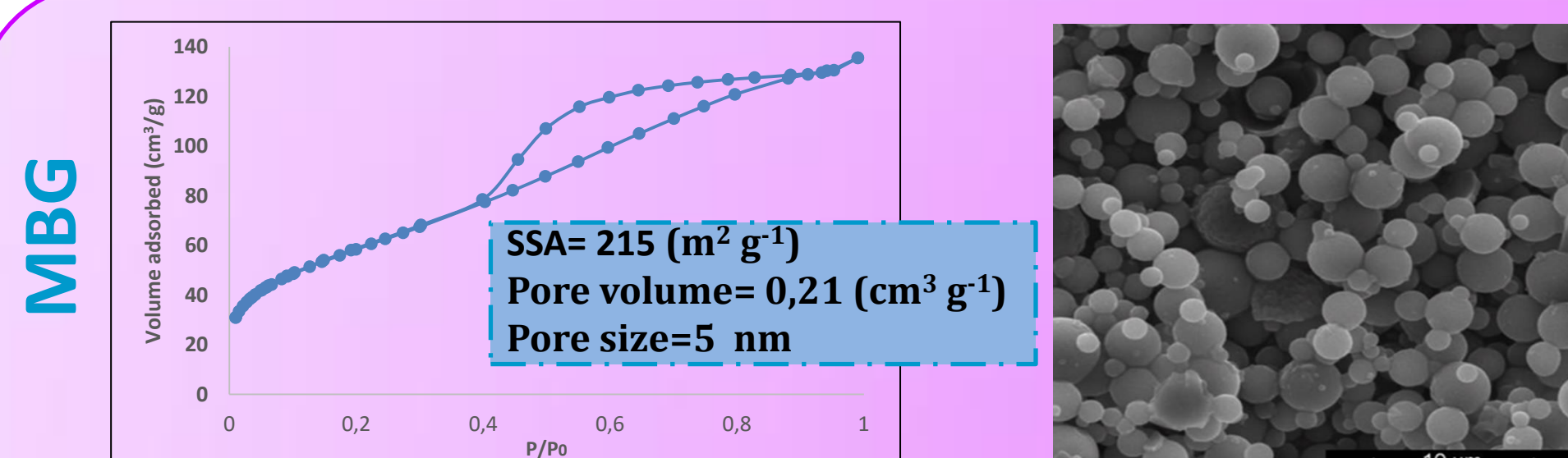
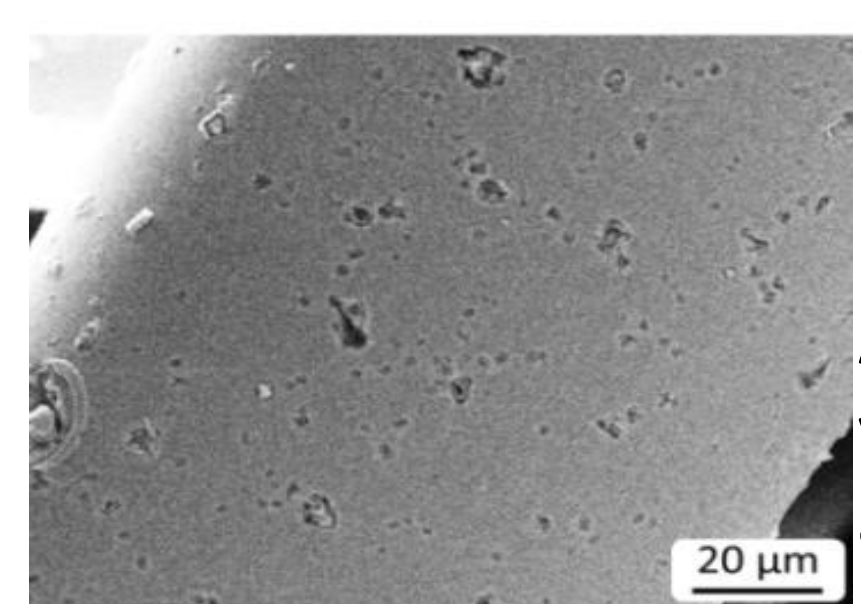


CEL2 are not well incorporated on fibre surface.



### Bioactivity test in SBF

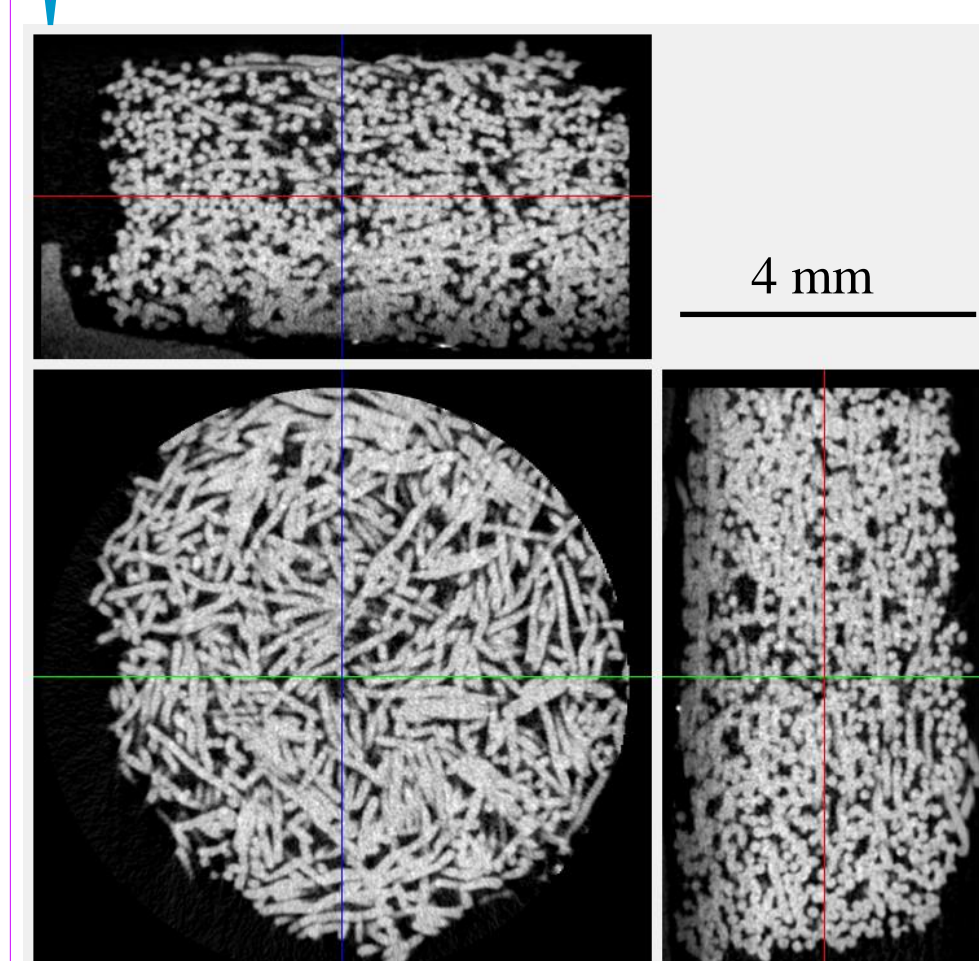
Only few particles still anchored to the scaffold



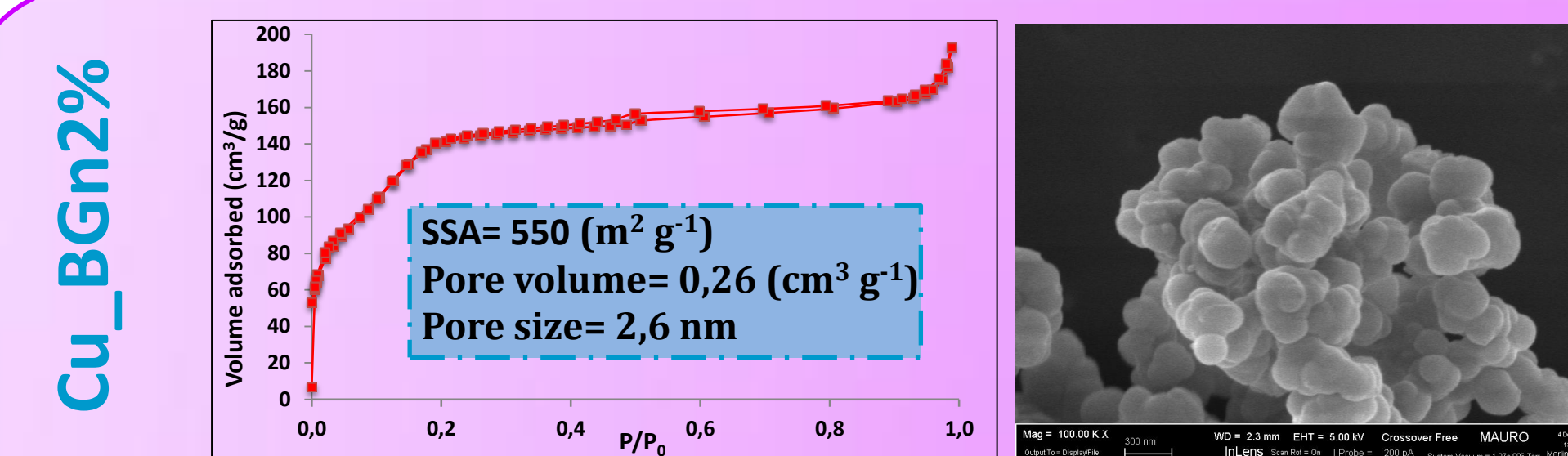
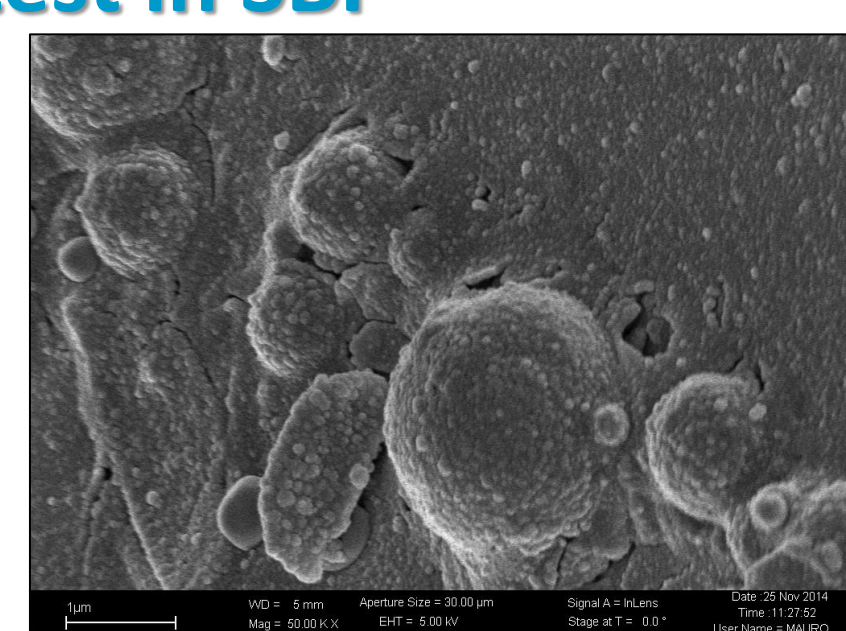
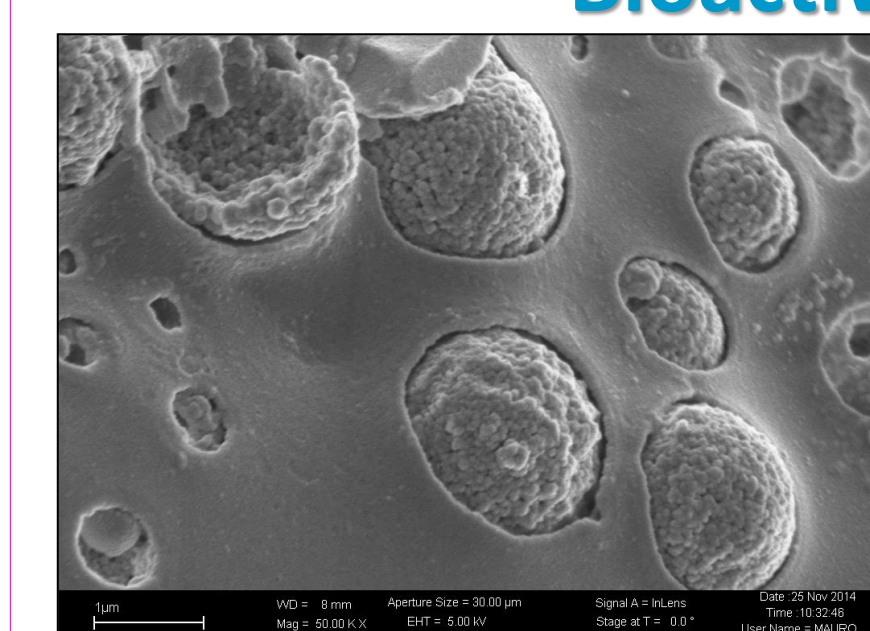
### S110\_MBG

- 5 wt.% bioactive powder
- Fibres: 6 mm

Total porosity: 53%  
Mean tr. Thick :159 µm  
Mean tr. Sep.: 203 µm



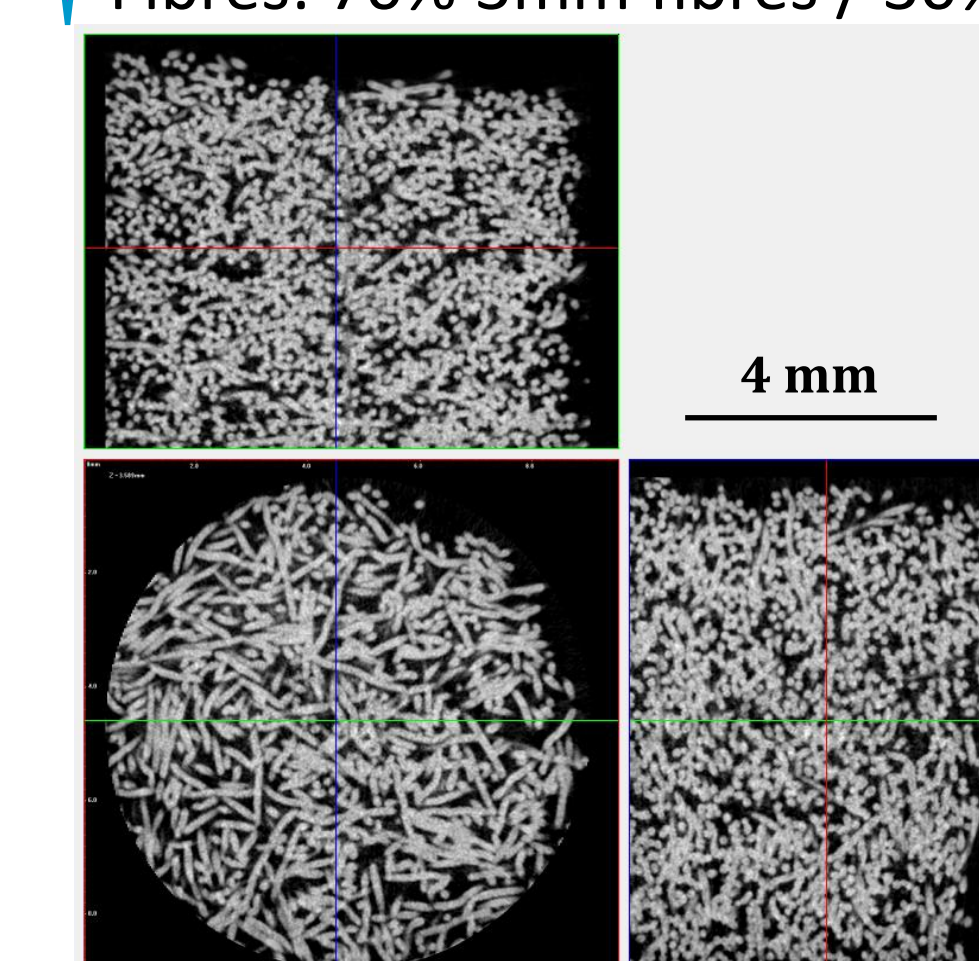
### Bioactivity test in SBF



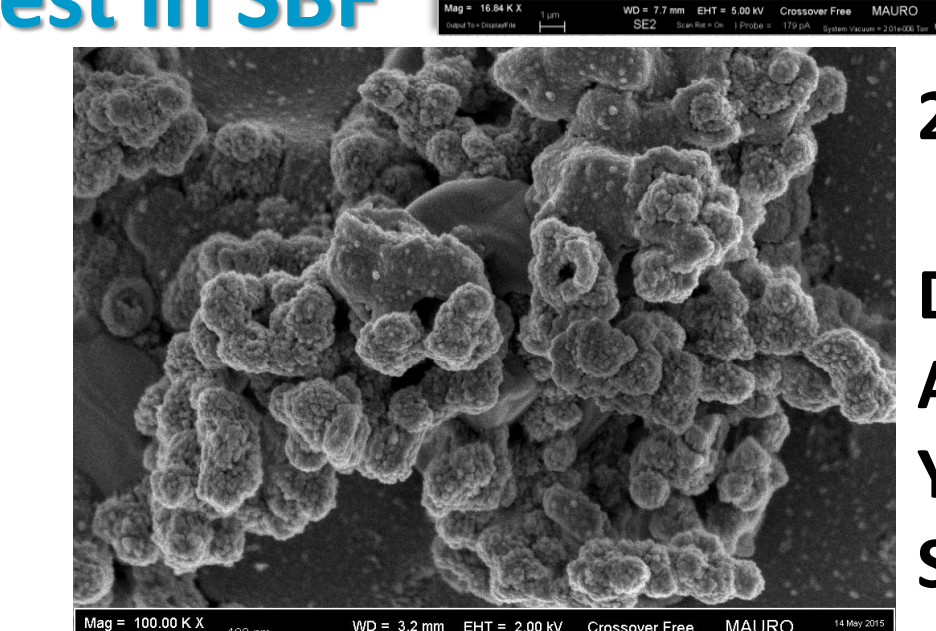
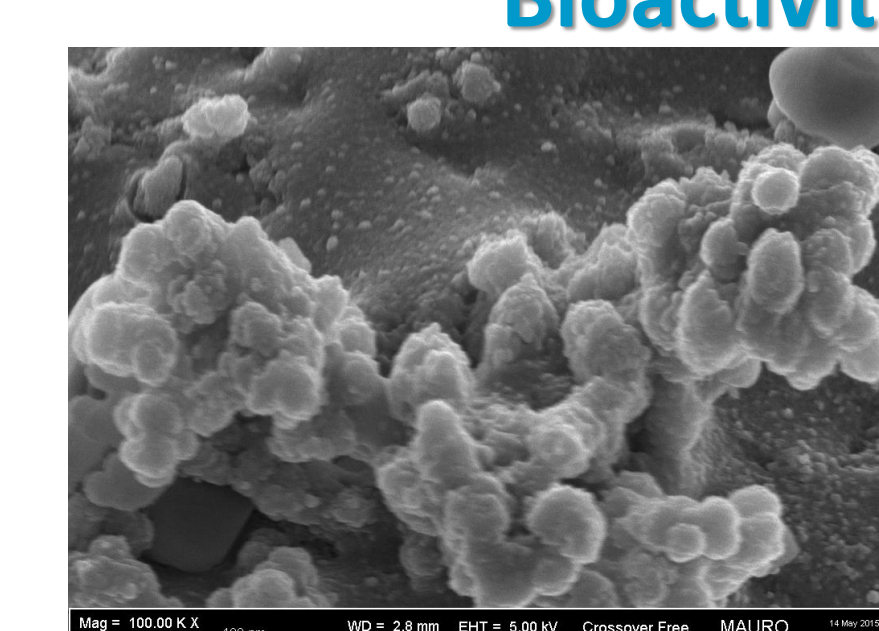
### S110\_Cu\_BGN2%

- 0.5 wt.% bioactive powder
- Fibres: 70% 3mm fibres / 30% 6mm fibres

Total porosity: 33%  
Mean tr. Thick :170 µm  
Mean tr. Sep.: 133 µm



### Bioactivity test in SBF



## Acknowledgement

The activity leading to this review has received funding from H2020-NMP-PILOTS-2015 under grant agreement no. 685872 (MOZART).

## References

- [1] D. Arcos et al. Chem Mater, 21 (2009), 1000–1009
- [2] C. Vitale-Brovarone et al. Mat Sci Eng C 31 (2) (2011) 434–442
- [3] Vitale Brovarone et al. Acta Biomater 3 (2007) 199–208
- [4] Vitale Brovarone et al. Key Engineering Materials Vol. 631 (2015) pp 43-47

## Conclusion

The incorporation of **MBG** and **Cu\_BGN2% powder** in the **phosphate glass fibrous scaffold** resulted to be a very interesting strategy to induce hydroxyapatite formation on the scaffold. Their fast bioactive response is due to their **mesoporosity**: it involves a high surface area available for ion exchange which is responsible for the glass bioactivity

These promising results encourage further investigation in order to fully exploit the ability of mesoporous particles to act as a system for **smart release of therapeutic ions and drugs**.